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<b>(54) Title:</b> TARGETED HETERO-ASSOCIATION OF RECOMBINANT PROTEINS TO MULTI-FUNCTIONAL COMPLEXES  <b>(57) Abstract</b>  The present invention relates to a method for targeted assembly of distinct active peptide or protein domains into a single complex and to such complexes. The invention relates particularly to the fusion of peptide or protein domains to complementary association domains which are derived from a single tertiary or quaternary structure by segmentation. The association domains are designed to assemble in a complementary fashion, thereby providing multifunctional (poly)peptides.		

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## TARGETED HETERO-ASSOCIATION OF RECOMBINANT PROTEINS TO MULTI-FUNCTIONAL COMPLEXES

### Background of the Invention

Increasingly, there is a need for proteins which combine two or more functions, such as binding or catalysis, in a single structure. Typically, proteins which combine two or more functions are prepared either as fusion proteins or through chemical conjugation of the component functional domains. Both of these approaches suffer from disadvantages. Genetic "single chain" fusions suffer the disadvantages that (i) only a few (2-3) proteins can be fused (Rock et al., 1992, *Prot. Eng.* 5, 583-591), (ii) mutual interference between the component domains may hinder folding, and (iii) the size of the fusion protein may make it difficult to prepare. The alternative, chemical cross-linking *in vitro* following purification of independently expressed proteins, is difficult to control and invariably leads to undefined products and to a severe loss in yield of functional material.

Recently, methods for achieving non-covalent association of two or more of the same functional domains have been developed. This can be achieved through the use of domains attached to peptides which self-associate to form homo-multimers (Pack & Plückthun, 1992, *Biochemistry* 31, 1579-1584). For example, the association of two separately expressed scFv antibody fragments by C-terminally fused amphipathic helices *in vivo* provides homo-dimers of antibody fragments in *E. coli* (PCT/EP93/00082; Pack et al., 1993, *Bio/Technology* 11, 1271-1277) or homo-tetramers; (Pack et al., 1995, *J. Mol. Biol.*, 246, 28-34).

To assemble distinct protein functions such as two antibody fragments with different specificities fused to such association domains, the helices must have a tendency to form hetero-multimers. In principle, this could be achieved with complementary helices such as the hetero-dimerizing JUN and FOS zippers of the AP-1 transcription factor (O'Shea et al., 1992, *Cell* 68, 699-708). The clear disadvantage of association domains based on hetero-associating helices,

however, is their pseudo-symmetry and their similar periodicity of hydrophobic and hydrophilic residues. This structural similarity results in a strong tendency to form homo-dimers and, thus, to lower significantly the yield of hetero-dimers (O'Shea et al., 1992, *Cell* 68, 699-708; Pack, 1994, Ph. D. thesis, Ludwig-Maximilians-Universität München). Furthermore, the formation of JUN/FOS hetero-dimers is kinetically disfavoured and requires a temperature-dependent unfolding of the kinetically favoured homo-dimers, especially JUN/JUN homo-dimers (PCT/EP93/00082; O'Shea et al., 1992, *Cell* 68, 699-708; Pack, 1994, Ph. D. thesis, Ludwig-Maximilians-Universität München). Because of the need for additional purification steps to separate the unwanted homo-dimers from hetero-dimers and the resulting decrease in yield, hetero-association domains based on amphipathic helices do not result in practical advantages compared to conventional chemical coupling.

These disadvantages of the prior art are overcome by the present invention which provides multi-functional polypeptides and methods for the preparation of these multi-functional proteins. This is achieved via the use of association domains which are designed to associate predominantly in a complementary fashion, and not to self-associate.

### Detailed Description of the Invention

In the earliest steps of protein folding, peptide chains form a disordered hydrophobic core by collapsing hydrophobic residues into the interior of an intermediate "molten globule". This hydrophobic effect is considered to be the most important driving force of folding (Matthews, 1993, *Annu. Rev. Biochem.* 62, 653 - 683; Fersht, 1993, *FEBS Letters* 325, 5 - 16). The burial of hydrophobic residues and the resulting exclusion of solvent is the determining factor in the stability of compact tertiary structures such as acyl-phosphatase (Pastore et al, *J. Mol. Biol.* 224, 427-440, 1992) interleukin-2 (Brandhuber et al., 1987, *Science* 238, 1707 - 1709), calbindin (Parmentier, 1990, *Adv. Exp. Med. Biol.* 269, 27-34) or ubiquitin (Briggs & Roder, 1992, *Proc. Natl. Acad. Sci. USA* 89, 2017 - 2021).

This concept forms the basis of the present invention, which provides individually encoded peptides or "segments" which, in a single continuous chain, would comprise a compact tertiary structure with a highly hydrophobic core. The component peptides are chosen so as to be asymmetric in their assumed structure, so as not to self-associate to form homo-multimers, but rather to associate in a complementary fashion, adopting a stable complex which resembles the parent tertiary structure. On the genetic level, these segments are encoded by interchangeable cassettes with suitable restriction sites. These standardized cassettes are fused C- or N-terminally to different recombinant proteins via a linker or hinge in a suitable expression vector system.

Thus, the present invention relates to a multi-functional polypeptide comprising:

- (a) a first amino acid sequence attached to at least one functional domain;
- (b) a second amino acid sequence attached to at least one further functional domain; and
- (c) optionally, further amino acid sequences each attached to at least one further functional domain;

wherein any one or more of said amino acid sequences interacts with at least one of said amino acid sequences in a complementary fashion to form a parental, native-like tertiary or optionally quaternary structure and wherein the parental, native-like tertiary or optionally quaternary structure is derived from a single parent polypeptide. In this context, the term parent polypeptide refers to a polypeptide which has a compact tertiary or quaternary structure with a hydrophobic core. The invention provides for many different parent polypeptides to be used as the basis for the association domain. Suitable polypeptides can be identified by searching for compact, single-domain proteins or protein fragments in the database of known protein structures (Protein Data Bank, PDB) and selecting structures that are stable and can be expressed at high yields in recombinant form. These structures can then be analyzed for hydrophobic sub-clusters by the method of Karpeisky and Ilyn (1992, *J. Mol. Biol.* 224, 629-638) or for structural units (such as  $\beta$ -elements or helical hairpin structures) by standard molecular modelling techniques. In a further embodiment, the present invention

provides for multi-functional polypeptides wherein the single parent polypeptide is taken from the list ubiquitin, acyl phosphatase, IL-2, calbindin and myoglobin.

In a preferred embodiment, the present invention provides a multi-functional polypeptide comprising two or more amino acid sequences each attached to at least one functional domain, wherein any two or more of said amino acid sequences can associate in a complementary fashion to provide a parental, native like, tertiary or optionally quaternary structure.

Once structural sub-domains are identified, the protein is dissected in such a way these sub-domains remain intact. The selection process can be expanded to proteins for which no structure is available but which satisfy the criteria of stability and good expression. For these proteins, folding sub-domains can be determined by hydrogen exchange pulse-labelling of backbone amides during the folding reaction, followed by NMR detection in the native state (Roder et al., 1988, *Nature* 355, 700-704; Udgaonkar & Baldwin, 1988, *Science* 255, 594-597). Alternatively, folding sub-domains can be identified by mild proteolysis, denaturation, purification of fragments and reconstitution *in vitro* (Tasayco & Carey, 1992, *Science* 255, 594-597; Wu et al., 1993, *Biochemistry* 32, 10271-10276). Finally, additional clues for the choice of cleavage sites can be obtained from the exon structure in the case of eukaryotic proteins, since the exons frequently (though not always) correspond to structural sub-domains of a protein. This has, for example, been discussed for the case of myoglobin (Go 1981, *Nature* 291, 90).

The yield of properly assembled molecules is expected to decrease significantly for constructs in which a protein domain is divided into three or more parts. This is due to the fact that several sub-domains must come together simultaneously to form a viable structure. This effect is countered by dividing the polypeptide chain into sub-domains that represent folding units (identified by the methods described above). Thus, not only the final, assembled complex but also assembly intermediates will have the stability necessary to allow their accumulation in the

host during expression, resulting in a greatly improved kinetic behaviour of the system.

In solution, the isolated segments have little secondary structure and remain monomeric or form transient, non-specific and easily disrupted aggregates. Only upon mixing, either by separate expression and purification, or by co-expression, can the concerted folding of complementary segments provide the necessary intermediate interaction of residues (Matthews, 1993, *Annu. Rev. Biochem.* 62, 653 - 683) that results in the formation of a compact, native-like structure. This association, mainly driven by the burial of hydrophobic residues of all segments into a single hydrophobic core, leads to a targeted assembly of the N- or C-terminally fused proteins to a multi-functional complex *in vivo* or *in vitro*.

Optionally, the reconstituted native-like structure may also contribute an enzymatic or binding activity to increase the number of effector functions in the assembled complex. Accordingly, the present invention also provides a multi-functional polypeptide as described above, in which the native-like, tertiary or quaternary structure provides a biological activity. For example, when acyl phosphatase is used as the basis of the association domain, it is expected that the multi-functional polypeptide will retain some phosphatase activity.

The present invention provides for many different types of functional domains to be linked into the multi-functional polypeptide. Particularly preferred are cases in which one or more, preferably two, of said functional domains are fragments derived from molecules of the immunoglobulin superfamily. In particularly preferred embodiments, said fragments are antibody fragments. Also preferred are cases in which at least one of the functional domains possesses biological activity other than that associated with a fragment derived from a member of the immunoglobulin superfamily. By way of example, the present invention provides for the targeted assembly of enzymes, toxins, cytokines, peptide hormones, immunoglobulins, metal binding domains, soluble receptors, lectins, lipoproteins, purification tails and bioactive peptides to multi-functional complexes (Fig. 1) based on a modular system of expression vectors, restriction sites and "plug-in" gene cassettes coding for assembly segments, peptide linkers and functional domains (Fig.2).

If covalent linkage between the segments is necessary to prevent dissociation at low concentrations, cysteines can be introduced to form inter-segmental disulphide bridges between the amino acid sequences which comprise the association domain (Ecker et al., 1989, *J. Biol. Chem.* 264, 1887-1893; Pack & Plückthun, 1992, *Biochemistry* 31, 1579-1584). Accordingly, the present invention provides multi-functional polypeptides wherein the folding of the component amino acid sequences is stabilized by a covalent bond.

In order to provide some flexibility between the association domain and the appended functional domains, it may be desired to incorporate a linker peptide. Accordingly, the present invention provides for multi-functional polypeptides of the type described above wherein at least one of the functional domains is coupled to said amino acid sequence via a flexible peptide linker. By way of example, the flexible linker may be derived from the hinge region of an antibody.

The invention enables even more complex multi-functional polypeptides to be constructed via the attachment of at least one further (poly)peptide to one or more of said amino acid sequences. By way of example, the further (poly)peptide can be taken from the list enzymes, toxins, cytokines, peptide hormones, immunoglobulins, metal binding domains, soluble receptors, lectins, lipoproteins, purification tails, in particular peptides which are able to bind to an independent binding entity, bioactive peptides, preferably of 5 to 15 amino acid residues, metal binding proteins, DNA binding domains, transcription factors and growth factors.

For therapeutic purposes, it is often desirable that proteinaceous substances display the minimum possible immunogenicity. Accordingly, the present invention provides for multi-functional polypeptides as described above in which at least one of said amino acid sequences, functional domains, or further (poly)peptides is of human origin.

In addition to the peptides and proteins provided above, the present invention also provides for DNA sequences, vectors, preferably bicistronic vectors, vector cassettes, characterised in that they comprise a DNA sequence encoding an amino acid sequence and optionally at least one further (poly)peptide comprised in the multifunctional polypeptide of the invention, and additionally at least one,

preferably singular cloning sites for inserting the DNA encoding at least one further functional domain or that they comprise DNA sequences encoding the amino acid sequences, and optionally the further (poly)peptide(s) comprised in the multifunctional polypeptide of the invention and suitable restriction sites for the cloning of DNA sequences encoding the functional domains, such that upon expression of the DNA sequences after the insertion of the DNA sequences encoding the functional domains into said restriction sites, in a suitable host the multifunctional polypeptide of the invention is formed. In a preferred embodiment said vector cassette is characterised in that it comprises the inserted DNA sequence(s) encoding said functional domain(s) and host cells transformed with at least one vector or vector cassette of the invention which can be used for the preparation of said multi-functional polypeptides.

In a further preferred embodiment, said host cell is a mammalian, preferably human, yeast, insect, plant or bacterial, preferably *E. coli* cell.

The invention further provides for a method for the production of a multifunctional polypeptide of the invention, which comprises culturing the host cell of the invention in a suitable medium, and recovering said multifunctional polypeptide produced by said host cell.

In a further embodiment, the invention relates to a method for the production of a multifunctional polypeptide of the invention which comprises culturing at least two host cells of the invention in a suitable medium, said host cells each producing only one of said first and said second amino acid sequences attached to at least one further functional domain, recovering the amino acid sequences, mixing thereof under mildly denaturing conditions and allowing in vitro folding of the multifunctional polypeptide of the invention from said amino acid sequences.

In a particular preferred embodiment, said method is characterised in that the further amino acid sequences attached to at least one further functional domain

are/is produced by at least one further host cell not producing said first or second amino acid sequence.

In another particularly preferred embodiment of the invention, said method is characterised in that at least one further amino acid sequence attached to at least one further functional domain is produced by the host cell of the invention producing said first or second amino acid sequence.

In further preferred embodiments, the present invention provides for pharmaceutical and diagnostic compositions comprising the multi-functional polypeptides described above, said pharmaceutical compositions optionally comprising a pharmaceutically acceptable carrier. Finally, the invention provides for a kit comprising one or more vector cassettes useful in the preparation of said multi-functional polypeptides.

The invention is now illustrated by reference to the following examples, which are provided for the purposes of illustration only and are not intended to limit the scope of the invention.

### Example 1: Segmented human ubiquitin as an assembly device

Ubiquitin is a compact intracellular protein of only 76 residues (Fig. 3) and a molecular weight of 5 kDa. It shows the highest conservation among all known proteins and is involved in the degradation pathway of intracellular eukaryotic proteins by forming intermediate isopeptide bonds to its C-terminus and to Lys48 (Hershko & Ciechanover, 1992, *Ann. Rev. Biochem.* 61, 761-807).

To use ubiquitin as an assembly device, the unwanted function can be abolished by truncation of the last three C-terminal residues (--Arg-Gly-Gly), and the exchange of Lys48 to Arg, which prevents the formation of isopeptide bonds to this residue. The altered sequence is then divided in a loop at position Gly36, so that the hydrophobic core falls apart into two segments (called ALPHA and BETA). The synthetic nucleotide sequence of the segments (Fig 4, 5) carry appropriate restriction sites (MroI-HindIII) at the termini, so that the cassette encoding the segments can be easily ligated to a EcoRI-MroI cassette encoding the flexible linker (hinge of hulgG3; Fig. 6). The cassettes are inserted into the expression vector pIG3 (EcoRI-HindIII; Fig. 7) encoding the scFv fragment of the antibody McPC603 under the lac promoter/operator (Ge et al., 1995, in: *Antibody engineering: A practical approach*. IRL Press, New York, Borrebaeck ed., 229-261). Insertion of a second functional fragment (scFv fragment of the anti- $\beta$ -lactam antibody 2H10 with phoA signal sequence) linked to association segment BETA as an XbaI-HindIII DNA fragment (Fig. 8) results in a di-cistronic expression vector (Pack, 1994, Ph. D. thesis, Ludwig-Maximilians-Universität München). After induction with IPTG and translation, the signal sequences guide the antibody fragments fused to the assembly segments to the periplasm, where they assemble to a complex with a reconstituted native-like ubiquitin fold and two different antibody specificities. The complex, a bispecific immunoglobulin, can be recovered and purified by affinity chromatography of cell extract (Pack, 1994, Ph. D. thesis, Ludwig-Maximilians-Universität München).

**Example 2: Covalent linkage of the native-like tertiary structure of the assembly device by engineered disulphide bridges and combination of a C-terminal peptide linker with an in-frame restriction site.**

The conformational stability of undivided, native ubiquitin can be enhanced by introduction of disulfides at positions 4 and 66 without perturbation in the backbone (Ecker et al., 1989, *J. Biol. Chem.* 264, 1887-1893; Fig. 9). In the context of this invention, the engineering of disulfide bridges provides the covalent linkage of segments (Fig. 10, 11) after co-folding and assembly.

To raise the number of possible functional domains in the assembled complex, a C-terminal peptide can be fused to one or more of the segments of the assembly device. To fuse a functional domain like an enzyme, cytokine, antibody fragment, purification peptide or toxin to this linker, a restriction site, preferably unique, has to be introduced in-frame (Fig. 11). Gene synthesis, cloning, expression as well as recovery of the assembled, covalently linked complex is according to example 1.

**Example 3: Segmented human interleukin-2 (IL2) as an assembly device**

Human interleukin-2 (Brandhuber et al., 1987, *Science* 238, 1707 - 1709; Kuziel & Greene, 1991, in: *The Cytokine Handbook*. Academic Press, 84-100) is used as an assembly device by segmentation between position His79 and Lys 80 (Fig. 12). The device, encoded by MroI-Ascl-HindIII gene cassettes (Fig. 13, 14) combines the low immunogenicity of the plasmatic protein with a preferable effector function of the native-like cytokine structure and an inter-segmental cysteine bridge (Cys58-Cys105) after assembly. The combination of one or more antibody fragments against tumor antigens with additional cytokines like IL6 or

IL12 targets the multi-cytokine complex (Rock et al., 1992, *Prot. Eng.* 5, 583-591) directly to the tumour.

#### **Example 4 : Segmented human apomyoglobin as an assembly device with three segments**

To use more than two segments of a native structure as an assembly device, the hydrophobic interface between the segments has to be large enough to provide the sufficient hydrophobic interaction for non-covalent linkage. Myoglobin (Fig.15) is expressible in large amounts in *E. coli* (Guillemette et al., 1991, *Protein Eng.* 4, 585-592). Up to six functional domains can be assembled by a threefold segmented structure (Fig. 16, 17, 18), three at the N-termini and three at the C-termini of the segments. The presence of heme additionally stabilizes the native-like apomyoglobin fold and can be used as a switch to influence the association constant of the multi-functional complex.

#### **Example 5: Bioactive peptides as functional domains**

Certain peptides derived from amphipathic loop structures of LPS-binding proteins (Hoess et al., 1993, *EMBO J.* 12, 3351-3356) are able to neutralize endotoxin. This effect is enhanced by multivalent display of these short peptides (10-15 residues; Hoess, unpublished results). The present invention provides a method to express and assemble several of short peptides (Fig.19), fused to an assembly segment, in a multivalent complex or in combination with other functional domains. The peptides can be fused either to the N-or to the C-terminus (Fig. 20, 21) of the assembly domain via the peptide linkers.

**Example 6: A purification tail for IMAC as a functional domain**

Peptide tails consisting of histidines are able to coordinate metal ions. They are used for purification of native proteins in immobilized metal affinity chromatography (IMAC). Multivalent display of the purification tail considerably improves the maximum purity achievable by IMAC (Lindner et al., 1992, *Methods: a companion to methods in enzymology* 4, 41-56). One or more gene cassettes (Fig. 22) encoding a polyhistidine tail can be fused to the assembly segment to provide a simple and efficient purification method for multi-functional complexes.

**Example 7: The platelet aggregation inhibitor decorsin as a functional domain**

Decorsin, a 39 residue protein of the leech *Macrobdella decora* (Fig. 23), acts as a potent antagonist of the platelet glycoprotein IIb-IIIa (Seymour et al., 1990, *J. Biol. Chem.* 265, 10143-10147). The gene cassette encoding the decorsin can be fused C- or N-terminally to an association segment (Fig. 24, 25). In arterial thrombotic diseases, a multivalent decorsin complex combined with an anti-fibrin antibody fragment can act as a powerful antithrombotic agent.

## Claims

1. A multifunctional polypeptide comprising:
  - (a) a first amino acid sequence attached to at least one functional domain;
  - (b) a second amino acid sequence attached to at least one further functional domain; and
  - (c) optionally, further amino acid sequences each attached to at least one further functional domain;wherein any one or more of said amino acid sequences interacts with at least one of said amino acid sequences in a complementary fashion to form a parental, native-like tertiary or optionally quaternary structure and wherein said parental, native-like tertiary or optionally quaternary structure is derived from a single parent polypeptide.
2. The multifunctional polypeptide according to claim 1, wherein said single parent polypeptide is ubiquitin, acyl-phosphatase, IL2, calbindin or apomyoglobin.
3. The multifunctional polypeptide according to claim 1 or 2, wherein said parental, native-like tertiary or optionally quaternary structure is biologically active.
4. The multifunctional polypeptide according to any one of claims 1 to 3, wherein at least one of said functional domains is a fragment derived from a member of the immunoglobulin superfamily.
5. The multifunctional polypeptide according to claim 4, wherein two of said functional domains are fragments derived from members of the immunoglobulin superfamily.

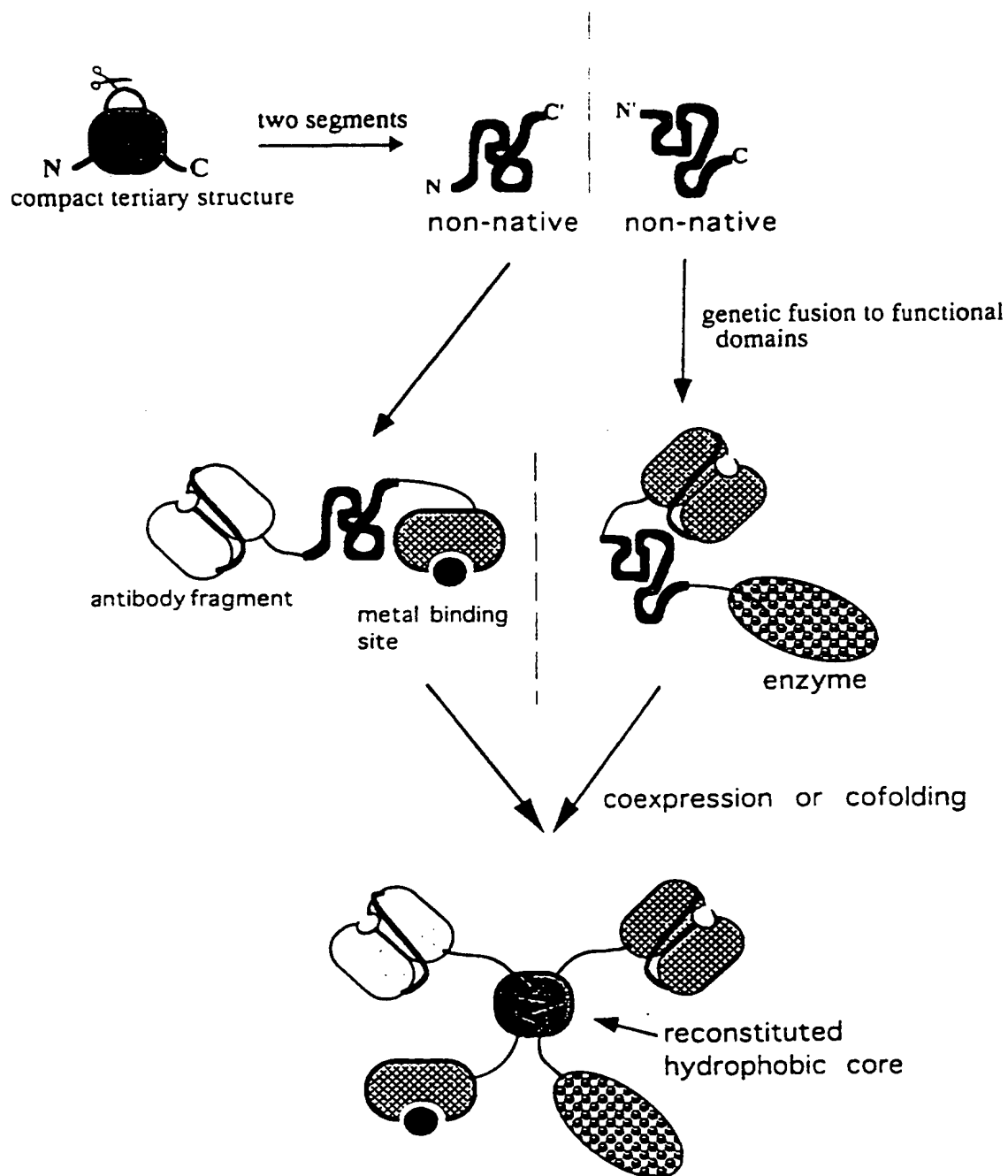
6. The multifunctional polypeptide according to claim 4 or 5, wherein said fragments are antibody fragments.
7. The multifunctional polypeptide according to any one of claims 1 to 6, wherein at least one of said functional domains is a biologically active molecule or a derivative thereof other than a fragment derived from a member of the immunoglobulin superfamily.
8. The multifunctional polypeptide according to any one of claims 1 to 6, wherein the folding of the amino acid sequences is stabilised by a covalent bonding.
9. The multifunctional polypeptide according to any one of claims 1 to 8, wherein at least one of said functional domains is coupled to said amino acid sequence(s) via a flexible peptide linker.
10. The multifunctional polypeptide according to claim 9, wherein said flexible peptide linker is an antibody hinge region.
11. The multifunctional polypeptide according to any one of claims 1 to 10, wherein at least one of said amino acid sequences is coupled to at least one further (poly)peptide.
12. The multifunctional polypeptide according to claim 11, wherein said further (poly)peptide is an enzyme, a toxin, a cytokine, a metal binding site, a metal binding protein, a soluble receptor, a DNA-binding domain, a transcription factor, an immunoglobulin, a bioactive peptide of 5 to 15 amino acid residues, a peptide hormone, a growth factor, a lectin, a lipoprotein, and a peptide which is able to bind to an independent binding entity.

13. The multifunctional polypeptide according to any one of claims 1 to 12, wherein at least one of said amino acid sequences, functional domains or further (poly)peptide(s) is of human origin.
14. A DNA sequence encoding an amino acid sequence and at least one functional domain and, optionally, at least one further functional (poly)peptide comprised in the multifunctional polypeptide of any one of claims 1 to 13.
15. A vector comprising at least one DNA molecule of claim 14.
16. The vector of claim 15, which is a bicistronic vector.
17. A vector cassette characterised in that it comprises a DNA sequence encoding an amino acid sequence and optionally at least one further (poly)peptide comprised in the multifunctional polypeptide of any one of claims 1 to 13, and additionally at least one, preferably a singular cloning site for inserting the DNA encoding at least one further functional domain.
18. A vector cassette characterised in that it comprises DNA sequences encoding the amino acid sequences, and optionally the further (poly)peptide(s) comprised in the multifunctional polypeptide of any one of claims 1 to 13, and suitable restriction sites for the cloning of DNA sequences encoding the functional domains, such that upon expression of the DNA sequences after the insertion of the DNA sequences encoding the functional domains into said restriction sites, in a suitable host the multifunctional polypeptide according to any one of claims 1 to 13 is formed.

19. The vector cassette according to claim 17 or 18 characterised in that it comprises the inserted DNA sequence(s) encoding said functional domain(s).
20. A host cell transformed with at least one vector according to claim 15 or 16, or at least one vector cassette according to claim 19.
21. The host cell according to claim 20, which is a mammalian, preferably human, yeast, insect, plant or bacterial, preferably E. coli cell.
22. A method for the production of a multifunctional polypeptide according to any one of claims 1 to 13, which comprises culturing the host cell according to claim 20 or 21 in a suitable medium, and recovering said multifunctional polypeptide produced by said host cell.
23. A method for the production of a multifunctional polypeptide according to any one of claims 1 to 13 which comprises culturing at least two host cells according to claim 20 or 21 in a suitable medium, said host cells each producing only one of said first and said second amino acid sequences attached to at least one further functional domain, recovering the amino acid sequences, mixing thereof under mildly denaturing conditions and allowing in vitro folding of the multifunctional polypeptide according to any one of claims 1 to 13 from said amino acid sequences.
24. The method according to claim 23, wherein the further amino acid sequence(s) (each) attached to at least one further functional domain are/is produced by at least one further host cell not producing said first or second amino acid sequence.

25. The method according to claim 23, wherein at least one further amino acid sequence attached to at least one further functional domain is produced by the host cell according to claim 20 or 21 producing said first or second amino acid sequence.
26. A pharmaceutical composition comprising the multifunctional polypeptide according to any one of claims 1 to 13 optionally in combination with a pharmaceutically acceptable carrier.
27. A diagnostic composition comprising the multifunctional polypeptide according to any one of claims 1 to 13.
28. A kit comprising at least one vector cassette according to claim 17 or 18.

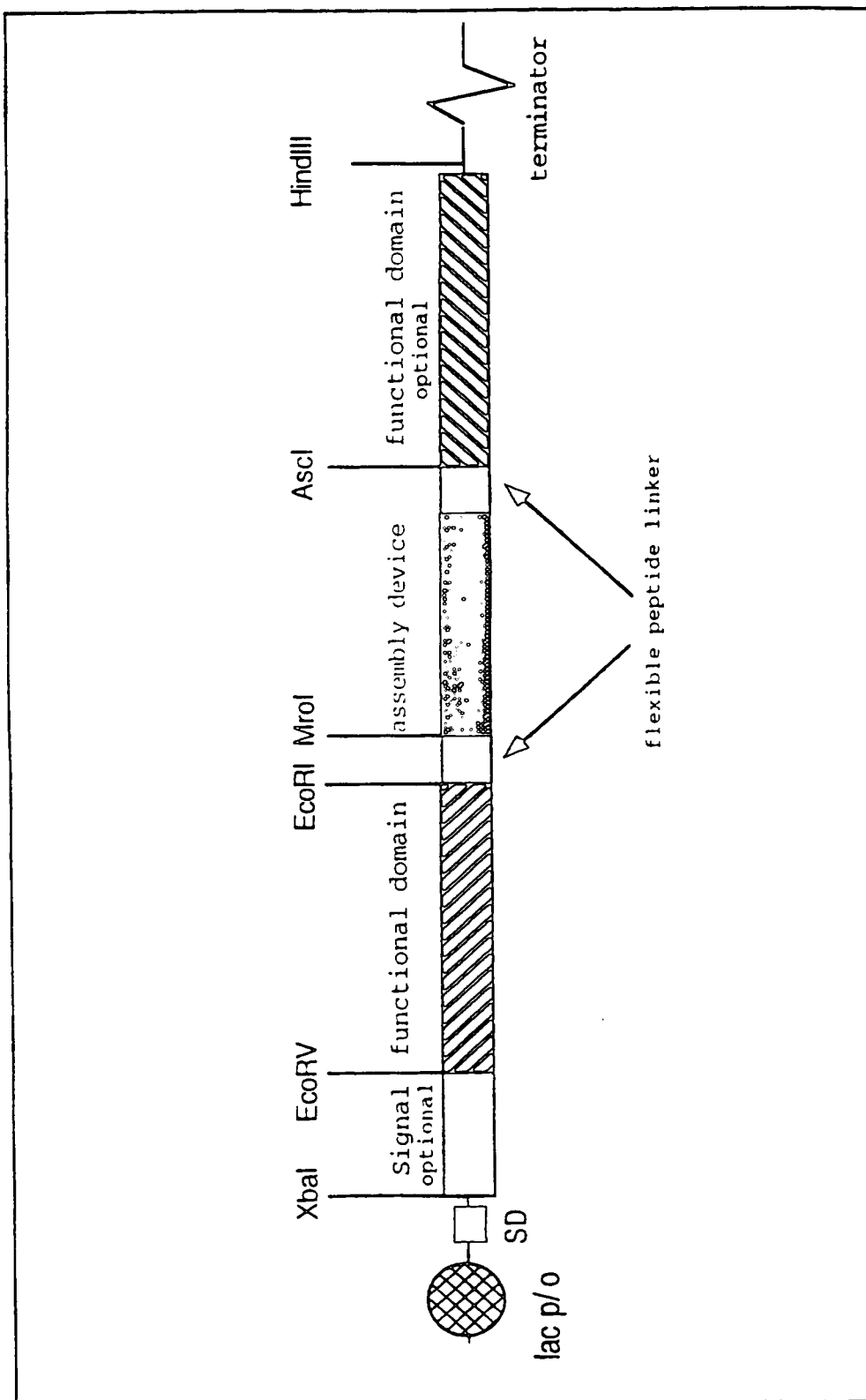
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**Fig.1:** segmented tertiary structure for a targeted hetero-association

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Fig. 2 : Modular cistron encoding functional domains  
N- and/or C-terminally fused to the assembly device



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Fig. 3 : protein sequence of human ubiquitin (segmented after Gly35)

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1      10      20      30      40      50
MQIFVKLTG KTITLEVEPS DTIENVKAKI QDKEGIPPDQ QRLIFAGKQL

      60      70
EDGRTLSDYN IQKESTLHLV LRLRGG**

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Fig. 4: MroI-Hind III gene cassette encoding for segment ALPHA of ubiquitin

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MroI
S  G  M  Q  I  F  V  K  T  L  T  G  K  T  I  T  L  E
TCC GGA ATG CAG ATC TTC GTT AAA ACC CTG ACC GGT AAA ACC ATC ACC CTG GAA
      9      18      27      36      45      54
AGG CCT TAC GTC TAG AAG CAA TTT TGG GAC TGG CCA TTT TGG TAG TGG GAC CTT

V  E  P  S  D  T  I  E  N  V  K  A  K  I  Q  D  K  E
GTT GAA CCG TCT GAC ACC ATC GAA AAC GTT AAA GCT AAA ATC CAG GAC AAA GAA
      63      72      81      90      99     108
CAA CTT GGC AGA CTG TGG TAG CTT TTG CAA TTT CGA TTT TAG GTC CTG TTT CTT

      HindIII
      G  *  *  A
GGT TGA TAA GCT T 3'
      117
CCA ACT ATT CGA A 5'

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Fig. 5: MroI-Hind III gene cassette encoding for segment BETA of ubiquitin

## MroI

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S   G   I   P   P   D   Q   Q   R   L   I   F   A   G   R   Q   L   E
TCC GGA ATC CCG CCG GAC CAG CAG CGT CTG ATC TTC GCT GGT CGT CAG CTG GAA
      9           18           27           36           45           54
AGG CCT TAG GGC GGC CTG GTC GTC GCA GAC TAG AAG CGA CCA GCA GTC GAC CTT

```

```

D   G   R   T   L   S   D   Y   N   I   Q   K   E   S   T   L   H   L
GAC GGT CGT ACC CTG TCT GAC TAC AAC ATC CAG AAA GAA TCT ACC CTG CAC CTG
      63           72           81           90           99           108
CTG CCA GCA TGG GAC AGA CTG ATG TTG TAG GTC TTT CTT AGA TGG GAC CTG GAC

```

## HindIII

```

V   L   R   L   *   *
GTT CTG CGT CTG TGA TAA 3'
      117           126
CAA GAC GCA GAC ACT ATT 5'

```

Fig. 6: EcoRI-MroI gene cassette encoding a flexible linker (huIgG3)

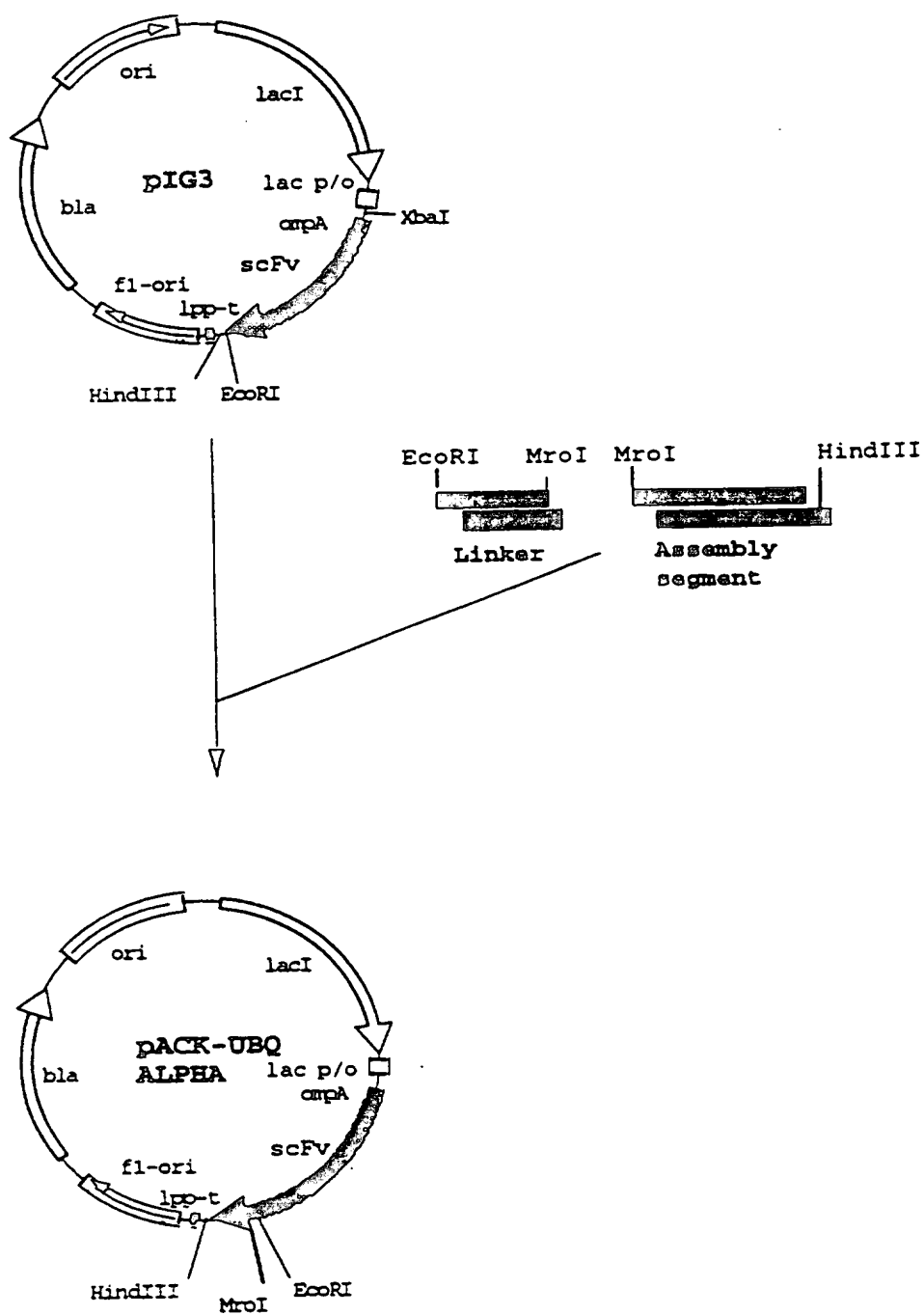
```

          EcoRI                               MroI
      E   F   T   P   L   G   D   T   T   H   T   S   G
5' GAA TTC ACC CCG CTG GGT GAC ACC ACC CAC ACC TCC GGA 3'
      9           18           27           36
3' CTT AAG TGG GGC GAC CCA CTG TGG TGG GTG TGG AGG CCT 5'

```

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Fig. 7 : Construction of monocistronic expression vector

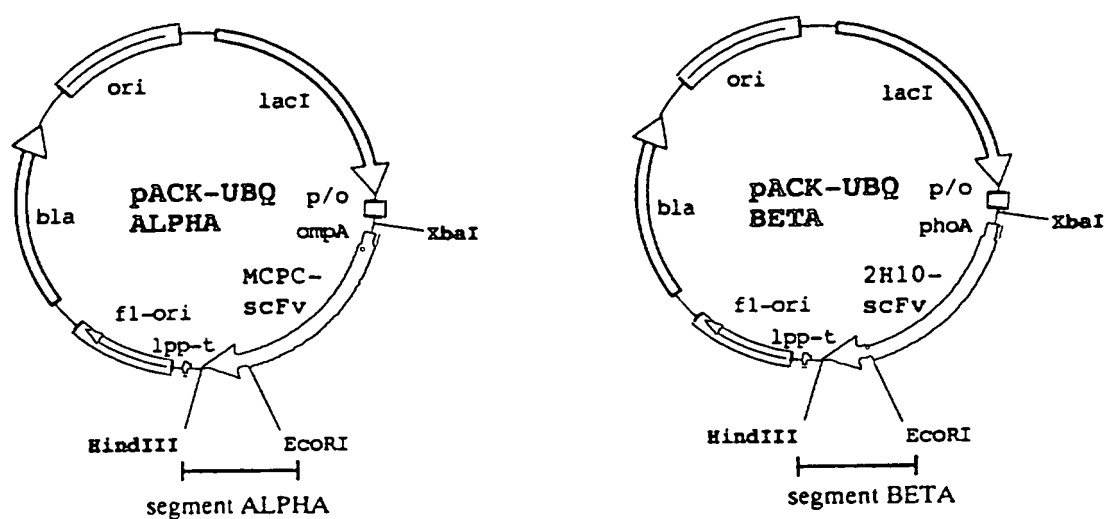


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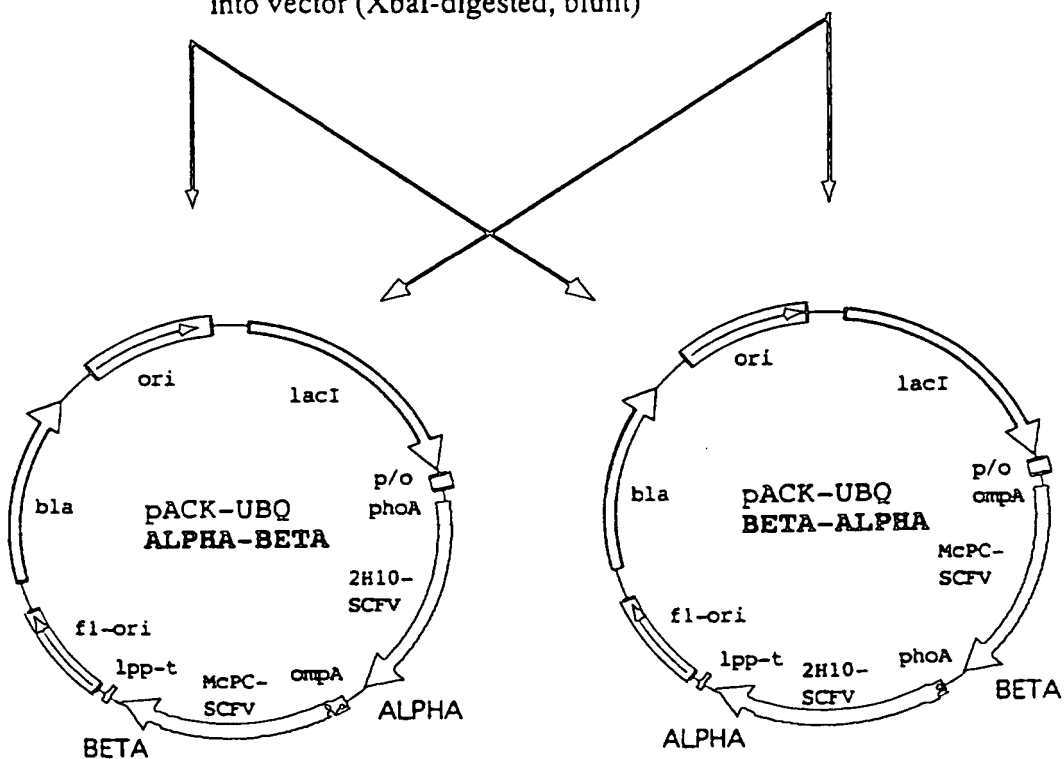
ISA/EP

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Fig. 8: Construction of dicistronic co-expression vectors



insert second cistron (XbaI/HindIII-digested, blunt)  
into vector (XbaI-digested, blunt)



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ISA/EP

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Fig. 9 : protein sequence of human ubiquitin with intersegmental disulfides Cys4 and Cys66 (segmented after Gly35)

```

1      10      20      30      40      50
MQICVKLTG KTITLEVEPS DTIENVKAKI QDKEGIPPDQ QRLIFAGKQL

      60      70
EDGRTLSDYN IQKESCLHLV LRLRGG**

```

Fig. 10: MroI-Hind III gene cassette encoding for segment ALPHA-CYS4 of ubiquitin

## MroI

```

S  G  M  Q  I  C  V  K  T  L  T  G  K  T  I  T  L  E
TCC GGA ATG CAG ATC TGC GTT AAA ACC CTG ACC GGT AAA ACC ATC ACC CTG GAA
      9      18      27      36      45      54
AGG CCT TAC GTC TAG ACG CAA TTT TGG GAC TGG CCA TTT TGG TAG TGG GAC CTT

V  E  P  S  D  T  I  E  N  V  K  A  K  I  Q  D  K  E
GTT GAA CCG TCT GAC ACC ATC GAA AAC GTT AAA GCT AAA ATC CAG GAC AAA GAA
      63      72      81      90      99      108
CAA CTT GGC AGA CTG TGG TAG CTT TTG CAA TTT CGA TTT TAG GTC CTG TTT CTT

```

## HindIII

```

G  *  *  A
GGT TGA TAA GCT T 3'
      117
CCA ACT ATT CGA A 5'

```

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Fig. 11 : MroI-AscI-Hind III gene cassette encoding for segment BETA-CYS66 with C-terminal GSGGAP linker of ubiquitin

## MroI

S G I P P D Q Q R L I F A G R Q L E  
 TCC GGA ATC CCG CCG GAC CAG CAG CGT CTG ATC TTC GCT GGT CGT CAG CTG GAA  
           9              18              27              36              45              54  
 AGG CCT TAG GGC GGC CTG GTC GTC GCA GAC TAG AAG CGA CCA GCA GTC GAC CTT

D G R T L S D Y N I Q K E S C L H L  
 GAC GGT CGT ACC CTG TCT GAC TAC AAC ATC CAG AAA GAA TCT TGC CTG CAC CTG  
           63              72              81              90              99              108  
 CTG CCA GCA TGG GAC AGA CTG ATG TTG TAG GTC TTT CTT AGA ACG GAC GTG GAC

                                  AscI                                  HindIII  
 V L R L G G S G G A P \* \*  
 GTT CTG CGT CTG GGG GGG AGC GGA GGC GCG CCG TGA TAA 3'  
           117              126  
 CAA GAC GCA GAC CCC CCC TCG CCT CCG CGC GGC ACT ATT 5'

Fig. 12: Protein sequence of human IL-2 (segmented after His79)

10          20          30          40          50          60  
 APTSSSTKKT QLQLEHLLLD LQMILNGINN YKNPKLTRML TFRFYMPKKA TELKHLQCLE  
 70          80          90          100          110          120  
 EELKPLEEVN NLAQSKNFHL RPRDLISNIN VIVLELKGSE TTFMCEYADE TATIVEFLNR  
 130  
 WITFCQSIIS TLT

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Fig.13 : MroI-AscI-Hind III gene cassette encoding for segment ALPHA of human IL-2

## MroI

S	G	A	P	T	S	S	S	T	K	K	T	Q	L	Q	L	E	H
TCC	GGA	GCA	CCT	ACT	TCA	AGT	TCT	ACA	AAG	AAA	ACA	CAG	CTA	CAA	CTG	GAG	CAT
		9			18			27			36			45			54
AGG	CCT	CGT	GGA	TGA	AGT	TCA	AGA	TGT	TTC	TTT	TGT	GTC	GAT	GTT	GAC	CTC	GTA

L	L	L	D	L	Q	M	I	L	N	G	I	N	N	Y	K	N	P
TTA	CTG	CTG	GAT	TTA	CAG	ATG	ATT	TTG	AAT	GGA	ATT	AAT	AAT	TAC	AAG	AAT	CCC
		63			72			81			90			99			108
AAT	GAC	GAC	CTA	AAT	GTC	TAC	TAA	AAC	TTA	CCT	TAA	TTA	TTA	ATG	TTC	TTA	GGG

K	L	T	R	M	L	T	F	K	F	Y	M	P	K	K	A	T	E
AAA	CTC	ACC	AGG	ATG	CTC	ACA	TTT	AAG	TTT	TAC	ATG	CCC	AAG	AAG	GCC	ACA	GAA
		117			126			135			144			153			162
TTT	GAG	TGG	TCC	TAC	GAG	TGT	AAA	TTC	AAA	ATG	TAC	GGG	TTC	TTC	CGG	TGT	CTT

L	K	H	L	Q	C	L	E	E	E	L	K	P	L	E	E	V	L
CTG	AAA	CAT	CTT	CAG	TGT	CTA	GAA	GAA	GAA	CTC	AAA	CCT	CTG	GAG	GAA	GTG	CTA
		171			180			189			198			207			216
GAC	TTT	GTA	GAA	GTC	ACA	GAT	CTT	CTT	CTT	GAG	TTT	GGA	GAC	CTC	CTT	CAC	GAT

														AscI		HindIII	
N	L	A	Q	S	K	N	F	H	G	G	S	G	G	A	P	*	
AAT	TTA	GCT	CAA	AGC	AAA	AAC	TTT	CAC	GGG	GGG	AGC	GGA	GGC	GCG	CCG	TGA	T
		225			234			243			252			261			
TTA	AAT	CGA	GTT	TCG	TTT	TTG	AAA	GTG	CCC	CCC	TCG	CCT	CCG	CGC	GGC	ACT	A

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Fig. 14 : MroI-AscI-Hind III gene cassette encoding for segment BETA of human IL-2

**MroI**

S	G	L	R	P	R	D	L	I	S	N	I	N	V	I	V	L	E
TCC	GGA	TTA	AGA	CCC	AGG	GAC	TTA	ATC	AGC	AAT	ATC	AAC	GTA	ATA	GTT	CTG	GAA
		9			18			27			36			45			54
AGG	CCT	AAT	TCT	GGG	TCC	CTG	AAT	TAG	TCG	TTA	TAG	TTG	CAT	TAT	CAA	GAC	CTT

L	K	G	S	E	T	T	F	M	C	E	Y	A	D	E	T	A	T
CTA	AAG	GGA	TCT	GAA	ACA	ACA	TTC	ATG	TGT	GAA	TAT	GCT	GAT	GAG	ACA	GCA	ACC
		63			72			81			90			99			108
GAT	TTC	CCT	AGA	CTT	TGT	TGT	AAG	TAC	ACA	CTT	ATA	CGA	CTA	CTC	TGT	CGT	TGG

I	V	E	F	L	N	R	W	I	T	F	C	Q	S	I	I	S	T
ATT	GTA	GAA	TTT	CTG	AAC	AGA	TGG	ATT	ACC	TTT	TGT	CAA	AGC	ATC	ATC	TCA	ACA
		117			126			135			144			153			162
TAA	CAT	CTT	AAA	GAC	TTG	TCT	ACC	TAA	TGG	AAA	ACA	GTT	TCG	TAG	TAG	AGT	TGT

**AscI****HindIII**

L	T	G	G	S	G	G	A	P	*
CTG	ACT	GGG	GGG	AGC	GGA	GGC	GCG	CCG	TGA T 3'
		171			180			189	
GAC	TGA	CCC	CCC	TCG	CCT	CCG	CGC	GGC	ACT A 5'

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Fig. 15 Protein sequence of human apomyoglobin (cut after Lys47 and Lys98)

```

mglsdgewql vlnvwgkvea dipghgqevl irlfkghpet lekfdkfkhl
51
ksedemkase dlkkhgatvl talggilkkk ghheaeikpl aqshatk hki
101
pvkylefise ciiqvlqskh pgdfgadaeg amnkalelfr kdmasykel
151
gfqg

```

Fig. 16 : MroI-AscI-Hind III gene cassette encoding for segment ALPHA of human apomyoglobin

## MroI

S	G	M	G	L	S	D	G	E	W	Q	L	V	L	N	V	W	G
TCC	GGA	ATG	GGT	CTG	TCT	GAC	GGT	GAA	TGG	CAG	CTG	GTT	CTG	AAC	GTT	TGG	GGT
		9			18			27			36			45			54
AGG	CCT	TAC	CCA	GAC	AGA	CTG	CCA	CTT	ACC	GTC	GAC	CAA	GAC	TTG	CAA	ACC	CCA

K	V	E	A	D	I	P	G	H	G	Q	E	V	L	I	R	L	F
AAA	GTT	GAA	GCT	GAC	ATC	CCG	GGT	CAC	GGT	CAG	GAA	GTT	CTG	ATC	CGT	CTG	TTC
		63			72			81			90			99			108
TTT	CAA	CTT	CGA	CTG	TAG	GGC	CCA	GTG	CCA	GTC	CTT	CAA	GAC	TAG	GCA	GAC	AAG

K	G	H	P	E	T	L	E	K	F	D	K	F	K	G	G	S	G
AAA	GGT	CAC	CCG	GAA	ACC	CTG	GAA	AAA	TTC	GAC	AAA	TTC	AAA	GGG	GGG	AGC	GGA
		117			126			135			144			153			162
TTT	CCA	GTG	GGC	CTT	TGG	GAC	CTT	TTT	AAG	CTG	TTT	AAG	TTT	CCC	CCC	TCG	CCT

AscI	HindIII		
G	A	P	*
GGC	GCG	CCG	TGA T 3'
		171	
CCG	GCG	GGC	ACT A 5'

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Fig. 17: MroI-AscI-Hind III gene cassette encoding for segment BETA of human apomyoglobin

## MroI

S	G	H	L	K	S	E	D	E	M	K	A	S	E	D	L	K	K
TCC	GGA	CAC	CTG	AAA	TCT	GAA	GAC	GAA	ATG	AAA	GCA	TCT	GAA	GAC	CTG	AAA	AAA
		9			18			27			36			45			54
AGG	CCT	GTG	GAC	TTT	AGA	CTT	CTG	CTT	TAC	TTT	CGT	AGA	CTT	CTG	GAC	TTT	TTT

H	G	A	T	V	L	T	A	L	G	G	I	L	K	K	K	G	H
CAC	GGT	GCT	ACC	GTT	CTG	ACC	GCT	CTG	GGT	GGT	ATC	CTG	AAA	AAA	AAA	GGT	CAC
		63			72			81			90			99			108
GTG	CCA	CGA	TGG	CAA	GAC	TGG	CGA	GAC	CCA	CCA	TAG	GAC	TTT	TTT	TTT	CCA	GTG

H	E	A	E	I	K	P	L	A	Q	S	H	A	T	K	H	K	G
CAC	GAA	GCT	GAA	ATC	AAA	CCG	CTG	GCT	CAG	TCT	CAC	GCT	ACC	AAA	CAC	AAA	GGG
		117			126			135			144			153			162
GTG	CTT	CGA	CTT	TAG	TTT	GGC	GAC	CGA	GTC	AGA	GTG	CGA	TGG	TTT	GTG	TTT	CCC

## AscI

## HindIII

G	S	G	G	A	P	*
GGG	AGC	GGA	GGC	GCG	CCG	TGA T 3'
		171			180	
CCC	TCG	CCT	CCG	CGC	GGC	ACT A 5'

K	A	L	E	L	F	R	K	D	M	A	S	N	Y	K	E	L	G
AAA	GCT	CTG	GAA	CTG	TTC	CGT	AAA	GAC	ATG	GCT	TCT	AAC	TAC	AAA	GAA	CTG	GGT
		117			126			135			144			153			162
TTT	CGA	GAC	CTT	GAC	AAG	GCA	TTT	CTG	TAC	CGA	AGA	TTG	ATG	TTT	CTT	GAC	CCA
F	S	Q	F	Q	E	T	F	V	H	S	R	V	V	F	F	Q	T

F	Q	G	G	G	S	G	G	A	P	*	HindIII
TTC	CAG	GGT	GGG	GGG	ACC	GGA	GGC	CGC	CCG	TGA	T 3'
		171			180			189			
AAG	GTC	CCA	CCC	CCC	TCG	CCT	CCG	CGC	GGC	ACT	A 5'

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Fig. 19: Peptide sequence of an endotoxin-neutralizing peptide as a functional domain

1                      11  
RWKVRKSFFKL Q

Fig.20: N-terminal EcoRV-EcoRI cassette encoding an endotoxin-neutralizing peptide

EcoRV														EcoRI			
I	M	R	W	K	V	R	K	S	F	F	K	L	Q	E	F		
5'	ATC	ATG	CGT	TGG	AAA	GTT	CGT	AAA	TCT	TTC	TTC	AAA	CTG	CAG	GAA	TTC	3'
			9			18			27			36			45		
3'	TAG	TAC	GCA	ACC	TTT	CAA	GCA	TTT	AGA	AAG	AAG	TTT	GAC	GTC	CTT	AAG	5'

Fig.21: C-terminal AscI-HindIII cassette encoding an endotoxin-neutralizing peptide

AscI														HindIII			
A	P	R	W	K	V	R	K	S	F	F	K	L	Q	*	*		
5'	GCG	CCG	CGT	TGG	AAA	GTT	CGT	AAA	TCT	TTC	TTC	AAA	CTG	CAG	TGA	TAA	3'
			9			18			27			36			45		
3'	CGC	GGC	GCA	ACC	TTT	CAA	GCA	TTT	AGA	AAG	AAG	TTT	GAC	GTC	ACT	ATT	5'

Fig. 22 AscI-HINDIII Gene cassette encoding a purification tail for IMAC

AscI								Hind III			
A	P	H	H	H	H	H	H	*	*		
5'	GCG	CCG	CAC	CAC	CAC	CAC	CAC	TGA	TAA	3'	
			9			18		27			
3'	CGC	GGC	GTG	GTG	GTG	GTG	GTG	CAC	ACT	ATT	5'

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Fig. 23 Protein sequence of the platelet aggregation inhibitor decorsin as a functional domain

```

1          11          21          31
APRLPQCQGD DQEKCLCNKD ECPPGQCRFP RGDADPYCE

```

Fig. 24 N-terminal EcoRV-EcoRI cassette encoding the platelet aggregation inhibitor decorsin

**EcoRV**

```

D I A P R L P Q C Q G D D Q E K C L
GAT ATC GCT CCG CGT CTG CCG CAG TGC CAG GGT GAC GAC CAG GAA AAA TGC CTG
          9          18          27          36          45          54
CTA TAG CGA GGC GCA GAC GGC GTC ACG GTC CCA CTG CTG GTC CTT TTT ACG GAC

```

```

C N K D E C P P G Q C R F P R G D A
TGC AAC AAA GAC GAA TGC CCG CCG GGT CAG TGC CGT TTC CCG CGT GGT GAC GCT
          63          72          81          90          99          108
ACG TTG TTT CTG CTT ACG GGC GGC CCA GTC ACG GCA AAG GGC GCA CCA CTG CGA

```

**EcoRI**

```

D P Y C E F
GAC CCG TAC TGC GAA TTC 3'
          117          126
CTG GGC ATG ACG CTT AAG 5'

```

Fig. 25 C-terminal AscI-HindIII cassette encoding the platelet aggregation inhibitor decorsin

**AscI**

```

A P A P R L P Q C Q G D D Q E K C L
GCG CCG GCT CCG CGT CTG CCG CAG TGC CAG GGT GAC GAC CAG GAA AAA TGC CTG
          12          21          30          39          48          57
CGC GGC CGA GGC GCA GAC GGC GTC ACG GTC CCA CTG CTG GTC CTT TTT ACG GAC

C N K D E C P P G Q C R F P R G D A
TGC AAC AAA GAC GAA TGC CCG CCG GGT CAG TGC CGT TTC CCG CGT GGT GAC GCT
          66          75          84          93          102          111
ACG TTG TTT CTG CTT ACG GGC GGC CCA GTC ACG GCA AAG GGC GCA CCA CTG CGA

```

**HindIII**

```

D P Y C E * *
GAC CCG TAC TGC GAA TGA TAA 3'
          120          129
CTG GGC ATG ACG CTT ACT ATT 5'

```

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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification:</b> <b>C12N 15/11, C07K 14/00</b>	<b>A3</b>	<b>(11) International Publication Number:</b> <b>WO 96/13583</b> <b>(43) International Publication Date:</b> 9 May 1996 (09.05.96)
<b>(21) International Application Number:</b> PCT/EP95/04117 <b>(22) International Filing Date:</b> 20 October 1995 (20.10.95) <b>(30) Priority Data:</b> 94116558.1      20 October 1994 (20.10.94)      EP <b>(34) Countries for which the regional or international application was filed:</b> DE et al. <b>(71) Applicant (for all designated States except US):</b> MORPHOSYS GESELLSCHAFT FÜR PROTEINOPTIMIERUNG MBH [DE/DE]; Frankfurter Ring 193a, D-80807 Munich (DE). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> PACK, Peter [DE/DE]; Franz-Wolter-Strasse 4/III, D-81925 Oberföhring (DE). LU-PAS, Andrei [DE/DE]; Fasanenstrasse 68a, D-82008 Unterhaching (DE). <b>(74) Agent:</b> VOSSIUS & PARTNER; P.O. Box 86 07 67, D-81634 Munich (DE).		<b>(81) Designated States:</b> JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i>  <b>(86) Date of publication of the international search report:</b> 13 February 1997 (13.02.97)
<b>(54) Title:</b> TARGETED HETERO-ASSOCIATION OF RECOMBINANT PROTEINS TO MULTI-FUNCTIONAL COMPLEXES <b>(57) Abstract</b> <p>The present invention relates to a method for targeted assembly of distinct active peptide or protein domains into a single complex and to such complexes. The invention relates particularly to the fusion of peptide or protein domains to complementary association domains which are derived from a single tertiary or quaternary structure by segmentation. The association domains are designed to assemble in a complementary fashion, thereby providing multifunctional (poly)peptides.</p>		

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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 95/04117

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 C12N15/11 C07K14/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 404 097 (BEHRINGWERKE) 27 December 1990 see the whole document ---	1
A	WO,A,94 20140 (THE USA GOVERNMENT) 15 September 1994 see the whole document ---	1
A	WO,A,93 15210 (MERCK PATENTS) 5 August 1993 see the whole document ---	1
A	WO,A,93 11162 (PROTEIN DESIGN LABS.) 10 June 1993 see the whole document ---	1
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☒ Further documents are listed in the continuation of box C.

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NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax (+31-70) 340-3016

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
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A	<p>P PACK: "Dissertation zur Erlangung des Doktorgrades der Fakultät für Chemie und Pharmazie der Ludwig-Maximilians-Universität München: Mini-Antikörper. Bivalente, tetravalente und bispezifische Immunglobuline aus E. coli "</p> <p>22 April 1994 , MUNICH, GERMANY XP002000449 cited in the application see the whole document -----</p>	1
A	<p>WO,A,92 10209 (WISTAR INSTITUTE) 25 June 1992 see the whole document -----</p>	1

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/EP 95/04117

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:  
Remark: Due to the unclear wording of the main claims and the rather theroretical nature of most examples provided, a complete search was deemed impossible for economical reasons. The search has limited to the subject matter of claim 1 as further limited by claims 2 and 4.
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because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searches without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 95/04117

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